BACE MedChemExpress

Product Data Sheet

Inhibitors • Screening Libraries • Proteins

(S,R,S)-AHPC-PEG2-NH2 dihydrochloride

Cat. No.:	HY-103603B	
CAS No.:	2341796-76-5	«N J
Molecular Formula:	$C_{28}H_{43}Cl_{2}N_{5}O_{6}S$	s
Molecular Weight:	648.64	нсі нсі
Target:	E3 Ligase Ligand-Linker Conjugates	
Pathway:	PROTAC	
Storage:	4°C, sealed storage, away from moisture	ОН
	* In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)	

BIOLOGICAL ACTIVITY		
Description	(S,R,S)-AHPC-PEG2-NH2 dihydrochloride (VH032-PEG2-NH2 dihydrochloride) is a synthesized E3 ligase ligand-linker conjugate that incorporates the (S,R,S)-AHPC based VHL ligand and 2-unit PEG linker used in the synthesis of PROTACs ^[1] .	
IC ₅₀ & Target	VHL	
In Vitro	PROTACs contain two different ligands connected by a linker; one is a ligand for an E3 ubiquitin ligase and the other is for the target protein. PROTACs exploit the intracellular ubiquitin-proteasome system to selectively degrade target proteins ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	

REFERENCES

[1]. Chan KH, et al. Impact of Target Warhead and Linkage Vector on Inducing Protein Degradation: Comparison of Bromodomain and Extra-Terminal (BET) Degraders Derived from Triazolodiazepine (JQ1) and Tetrahydroquinoline (I-BET726) BET Inhibitor Scaffolds. J Me

Caution: Product has not been fully validated for medical applications. For research use only.

 Tel: 609-228-6898
 Fax: 609-228-5909
 E-mail: tech@MedChemExpress.com

 Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA